PA . _NT COOPERATION TREAT.

	From the INTERNATIONAL BUREAU				
PCT	To:				
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THE PROPERTY OF THE POTTON	United States Patent and Trademark				
NOTIFICATION OF ELECTION	Office				
(PCT Rule 61.2)	(Box PCT)				
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	Washington, DC 20231 ETATS-UNIS D'AMERIQUE				
	ETATS-UNIS D'AIMENIQUE				
Date of mailing (day/month/year)	in its capacity as elected Office				
04 August 1998 (04.08.98)					
International application No.	Applicant's or agent's file reference				
PCT/GB97/03400					
International filing date (day/month/year)	Priority date (day/month/year)				
10 December 1997 (10.12.97)	10 December 1996 (10.12.96)				
Applicant					
KING, David, John et al					
The designated Office is hereby notified of its election mad	e:				
X in the demand filed with the International Preliminary	/ Examining Authority on:				
08 July 1998 (08.07.98)				
in a notice effecting later election filed with the Intere	national Bureau on:				
2. The election X was					
was not					
made before the expiration of 19 months from the priority	date or, where Rule 32 applies, within the time limit under				
Rule 32.2(b).					
	Authorized officer				
The International Bureau of WIPO					
34, chemin des Colombettes 1211 Geneva 20, Switzerland	P. Regis				
Facsimile No.: (41-22) 740.14.35	Telephone No.: (41-22) 338.83.38				
FacSittile No.: (41-22) 740.14.00	0100570				

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INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

			(PC) Article 36 a	iliu inule	, 0,	
Applicant's or a	gent's	file reference	FOR FURTHER ACT	TION	See I Prelin	Notification of Transmittal of International minary Examination Report (PCT/IPEA/416)
18451WO	-li-otic	n No	International filing date (day/mo	onth/year)		Priority date (day/month/year)
International application No. International filing date (day/molecules) PCT/GB97/03400 10/12/1997					10/12/1996	
			ational classification and IPC			
	atent C	lassification (IFO) of the	adorial electrical			
C07K16/00						
Applicant	TUE	RAPEUTICS LIMI	TFD et al.			
JELLTECH	INC	HAP LOTICO LIVII	123 015			
and is tr	ansmi	tted to the applicant	according to Article 30.			ernational Preliminary Examining Authority
2. This RE	PORT	consists of a total of	of 4 sheets, including this co	over sheet.		
wh bef These a	ich ha fore th	ve been amended a is Authority (see Ru es consist of a total	of sheets.	the Admini	sheets	s containing rectifications made ve Instructions under the PCT).
3. This rep	oort co		elating to the following items:			
!	Ø	Basis of the report				
11		Priority	e of eninion with regard to no	veltv. inver	ntive s	step and industrial applicability
111						·
IV 		Lack of unity of inv	rention ant under Article 35(2) with re	egard to no	velty,	, inventive step or industrial applicability;
V	×	citations and expla	anations supporting such stat	tement	•	
VI		Certain documents				
VII	\boxtimes		the international application			·
VIII		Certain observation	ons on the international applic	cation		
Date of sub	missio	n of the demand		Date of com	pletio	n of this report
08/07/19						7 9. 01. 99
Name and	mailing	address of the IPEA/		Authorized	office	Spinister S Million
 	Eu	ropean Patent Office		Winchliff	P	

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D-80298 Munich

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/GB97/03400

I.	Basis	of the	report
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1. This report has been drawn on the basis of (substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments.):

-	ne n	aport since they d	
!	Desc	cription, pages:	
	1-28		as originally filed
	Claiı	ns, No.:	
	1-10		as originally filed
	Drav	wings, sheets:	
	1/14	-14/14	as originally filed
2.	The	amendments hav	ve resulted in the cancellation of:
		the description,	pages:
		the claims,	Nos.:
		the drawings,	sheets:
3.		This report has be	peen established as if (some of) the amendments had not been made, since they have been been been been been been been be
4	Ado	ditional observation	ons, if necessary:

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/GB97/03400

- V. Reasoned statement under Article 35(2) with r gard to novelty, inv ntive st p or industrial applicability; citations and explanations supporting such statement
- 1. Statement

Novelty (N)

Yes:

Yes:

Claims

No: Claims 1-10

Inventive step (IS)

Yes: Claims

No: Claims 1-10

Industrial applicability (IA)

Claims 1-10

No: Claims

2. Citations and explanations

see separate sheet

VII. Certain defects in the international application

The following defects in the form or contents of the international application have been noted:

see separate sheet

Item V

Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

- 1. The subject matter of claims 1 to 10 does not fulfill the requirements of Article 33(2) PCT because it is not novel.
- 1.1 British Journal of Cancer (1994), **70:1126-1130**, Pedley R. et al (D1) discloses in the materials and methods section a method of producing randomly modified Fab' which is identical to that found on p.16 of the description. It is assumed that the method disclosed in D1 allows attachment of PEG to cysteine residues on the Fab'. Consequently the subject matter of claims 1 to 10 is identical to that disclosed in D1.
- 1.2 WO 96 09325 (D2) discloses on p. 6, lines 20-38, a method to conjugate PEG to the thiol groups of antibodies. After PEGylation, the whole antibodies are proteolytically cleaved in order to produce Fab' conjugated to PEG. Consequently the subject matter of claims 1 to 10 is identical to that disclosed in D2.

Re Item VII

Certain defects in the international application

1. Contrary to the requirements of Rule 5.1(a)(ii) PCT, the relevant background art disclosed in the document D2 is not mentioned in the description, nor is this documents identified therein.

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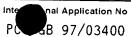
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INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference		of Transmittal of International Search Report 220) as well as, where applicable, item 5 below.
International application No.	International filing date (day/month/year)	(Earliest) Priority Date (day/month/year)
PCT/GB 97/03400	10/12/1997	10/12/1996
Applicant CELLTECH THERAPEUTICS L	IMITED et al.	
	been prepared by this International Searching Aut g transmitted to the International Bureau.	hority and is transmitted to the applicant
This International Search Report cons X It is also accompanied by a companied by	ists of a total of3 sheets. copy of each prior art document cited in this report	
Certain claims were found	unsearchable(see Box I).	
2. Unity of invention is lacking	ng(see Box II).	
	contains disclosure of a nucleotide and/or amin- ried out on the basis of the sequence listing	o acid sequence listing and the
	filed with the international application.	
	furnished by the applicant separately from the inte	rnational application,
,	but not accompanied by a statement to the matter going beyond the disclosure in the	
	Transcribed by this Authority	
4. With regard to the title, χ	the text is approved as submitted by the applicant	
	the text has been established by this Authority to re	ead as follows:
5. With regard to the abstract,	the tout is approved as submitted by the applicant	·
	the text is approved as submitted by the applicant the text has been established, according to Rule 3 Box III. The applicant may, within one month from	8.2(b), by this Authority as it appears in
	Search Report, submit comments to this Authority	
C. The figure of the description is the	whilehed with the chatract is	
6. The figure of the drawings to be p		None of the figures.
	as suggested by the applicant.	[A] Notice of the lightes.
	because the applicant failed to suggest a figure. because this figure better characterizes the inventi	ion
<u> </u>		···

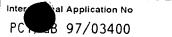
INTERNATIONAL SEARCH REPORT



IPC 6	C07K16/00		
According	to International Patent Classification(IPC) or to both national clas	sification and IPC	
	SEARCHED		
IPC 6	documentation searched (classification system followed by classif $C0.7K$	ication symbols)	
	ation searched other than minimumdocumentation to the extent th		
Electronic	data base consulted during the international search (name of data	a base and, where practical, sea	arch terms used)
C. DOCUM	ENTS CONSIDERED TO BE RELEVANT		
Category °	Citation of document, with indication, where appropriate, of the	relevant passages	Relevant to claim No.
X	PEDLEY R. B. ET AL.,: "The pot enhanced tumour localisation by poly(ethylene glycol) modificat anti-CEA antibody" BR. J. CANCER, vol. 70, - 1994	/	1-10
	pages 1126-1130, XP002061757 see the whole document		
Χ .	WO 96 09325 A (IMMUNOMEDICS INC 1996 see the whole document) 28 March	1-10
		-/	
X Furthe	er documents are listed in the continuation of box C.	χ Patent family memb	ers are listed in annex.
	egories of cited documents :	_ <u> </u>	after the international filing date
conside E" earlier do filing da		cited to understand the invention "X" document of particular re	In conflict with the application but principle or theory underlying the
citation	it which may throw doubts on priority claim(s) or cited to establish the publication date of another or other special reason (as specified) nt referring to an oral disclosure, use, exhibition or	"Y" document of particular re cannot be considered to	ovel or cannot be considered to p when the document is taken alone levance; the claimed invention involve an inventive step when the
oiner me P" documen"	eans It published prior to the international filing date but In the priority date claimed	ments, such combinatio in the art. "&" document member of the	with one or more other such docu- n being obvious to a person skilled
Date of the ac	ctual completion of theinternational search	Date of mailing of the inte	
. 8	April 1998	24/04/1998	·
Name and ma	illing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk	Authorized officer	
	Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Müller, F	

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INTERNATIONAL SEARCH REPORT



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C.(Continue Category °	ation) DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with indication, where appropriate, of the relevant passages		Polovent to elsies No		
J - ',	The state of the s		Relevant to claim No.		
A	CHIEN-TSUN KUAN ET AL: "PSEUDOMONAS EXTOXIN A MUTANTS REPLACEMENT OF SURFACE EXPOSED RESIDUES IN DOMAIN IN WITH CYSTEINERESIDUES THAT CAN BE MODIFIED WITH POLYETHYLENE GLYCOL IN A SITE-SPECIFIC MANNER" JOURNAL OF BIOLOGICAL CHEMISTRY, vol. 269, no. 10, 11 March 1994, pages 7610-7616, XP000563744 see the whole document		1-10		
4	WOGHIREN C. ET AL.,: "Protected Thiol-Polyethylen glycol: a new activated polymer for reversible protein modification" BIOCONJUGATE CHEM., vol. 4, - 1993 pages 314-318, XP002061758		1-10		
	see the whole document				
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INTERNATIONAL SEARCH REPORT

Infor

on patent family members

Integral Application No
PC - GB 97/03400

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9609325 A	28-03-96	US 5670132 A AU 3674995 A CA 2200482 A EP 0784634 A	23-09-97 09-04-96 28-03-96 23-07-97

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INTERNATIONAL APPLICATION PUBLIS		UNDER THE PATENT COOPERATION	N TREATY (PCT)
(51) Internati nal Patent Classification 6 : C07K 16/00	Al	(11) International Publication Number: (43) International Publication Date	WO 98/25971
(21) International Application Number: PCT/Gg (22) International Filing Date: 10 December 1997 (BY, CA, CH, CN, CU, CZ, DE, GH, GM, HU, ID, IL, IS, IP, K LK, LK, LS, LT, LU, LV, MD	DK, EE, ES, FI, GB, GE, E. KG, KP, KR, K2, I.C. MG MK MN MW MX
(30) Priority Data: 9625640.9 (0 December 1996 (10.12.9	6) G	NO, NZ, PL, PT, RO, RU, SD, SI TR, TT, UA, UG, US, UZ, VN (CH, GM, KE, LS, MW, SD, SZ, (AM, AZ, BY, KG, KZ, MD, RU	E, SG, SI, SK, SL, TI, TM, YU, ZW, ARIPO patent UG, ZW), Eurusian patent TI, TM), Eurusian patent
(71) Applicant for all designated States except US): CE THERAPEUTICS LIMITED [GB/GB], 216 Bu Stough, Berkshire SL1 4EN (GB)	LLTEC th Row	(AT. BE, CH. DE, DK, ES, Ft. MC, NL, PT, SE), OAPI patent (GA, CN, MI., MR, NE, SN, TD)	BE BLOE CO OLOM

(72) Inventors; and

- (75) Inventor Applicants (for US only): KING. David, John [GB/GB] 69 Watchetts Drive. Camberley, Surrey GU15 2PF (GB). CHAPMAN. Andrew. Paul [GB/GB]; 48 Hanworth Road, Hampton, Middlesex TW12 3DL (GB).
- (74) Agent: MERCER, Christopher, Paul, Carpmaels & Ransford, 43 Bloomsbury Square, London WC1A 2RA (GB)

Published

With international search report.

Before the expiration of the time that for amending the claims and to be republished in the event of the receipt of amendments.

(54) Title: MONOVALENT ANTIBODY FRAGMENTS

(57) Abstract

Monovalent antibody fragments are described, each of which has one or more polymer molecules site-specifically attached through a sulphur atom of a cysteme residue located outside of the variable region domain of the antibody. The polymers include synthetic of naturally occurring polymers such as polyalkylenes, polyalkylenes, polyaxyalkylenes or polysaccharides. Each fragment may be attached to one or more effector or reporter molecules, and is of use in therapy or diagnostics where it has markedly improved binding and/or pharmacokinetic properties when compared to other antibody fragments which have the same number and type of polymer molecules, but in which the polymer molecules are randomly attached.

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CLAIMS

- A modified monovalent antibody fragment comprising a monovalent antibody fragment and at least one polymer molecule in covalent linkage characterised in that each cysteine residue located in the antibody fragment outside of the variable region domain of the fragment is either covalently linked through its sulphur atom to a polymer molecule or is in disulphide linkage with a second cysteine residue located in the fragment provided that at least one of said cysteine residues is linked to a polymer molecule.
- An antibody fragment according to Claim 1 which is covalently linked to one, two or three polymer molecules through one, two or three cysteine residues located in the fragment outside of its variable region domain.
- An antibody fragment according to Claim 1 or Claim 2 wherein the polymer is an optionally substituted straight or branched chain polyalkylene, polyalkenylene or polyoxyalkylene polymer or a branched or unbranched polysaccharide.
- An antibody fragment according to Claim 3 wherein the polymer is an optionally substituted straight or branched chain poly(ethylene glycol), poly(propylene glycol) or poly(vinyl alcohol) and derivatives thereof.
 - 5. An antibody fragment according to Claim 4 wherein the polymer is methoxy(polyethylene glycol) and derivatives thereof.
 - 6. An antibody fragment according to any one of Claim 1 to Claim 5 in which the variable region domain is monomeric and comprises an immunoglobulin heavy (VH) or light (VL) chain variable domain, or is dimeric and contains VHVH, VHVL or VL-VL dimers in which the VH and VL chains are non-covalently associated or covalently coupled.

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- 7. An antibody fragment acc rding to Claim 6 wherein each V_H and/ r V_L domain is covalently attached at a C-t minal amino acid to at least on other antibody domain r a fragment thereof.
- An antibody fragment according to Claim 7 which is a Fab or Fab' fragment.
 - An antibody fragment according to any one of Claim 1 to Claim 8
 covalently attached to one or more effector or reporter molecules.
 - 10. A pharmaceutical composition comprising a monovalent antibody fragment according to any of the preceding claims together with one or more pharamaceutically acceptable excipients, diluents or carriers.